Palisaded and Neutrophilic Granulomatous Dermatitis in a Patient with Dermatomyositis and Nasopharyngeal Cancer

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Abstract

Palisaded and Neutrophilic Granulomatous Dermatitis (PNGD) is a rare form of neutrophilic dermatoses with granulomatous inflammation. It is a benign, inflammatory dermatosis with distinct histopathological features and varied clinical presentations. Associated systemic conditions include connective-tissue diseases and lymphoproliferative disorders. Dermatomyositis associated PNGD was rarely reported before. Herein, a PNGD patient with dermatomyositis and nasopharyngeal cancer was presented. The clinical and pathological findings as well as mechanism of neutrophilic and granulomatous infiltrates in autoimmune connective tissue disease related skin lesions were also briefly discussed.

Keywords: Palisaded and Dermatitis; Dermatomyositis; Neutrophilic Granulomatous; Nasopharyngeal Cancer;

Case Report

A 42-year-old male patient went to our clinic in 2004, presented with erythematous patches and plaques on nasal base, nose and bilateral hands for 4 months. There was no oral ulcer, muscular weakness, or arthralgia. The clinical impression was tumid lupus erythematosus. The laboratory examination results revealed a positive antinuclear antibody (ANA: 1:1280, speckled type), positive anti-La antibody, and negative anti-Ro, anti-Sm/RNP, anti-Scl 70 antibodies. The complement C3 and C4, complete blood count, liver and renal profiles were all within normal limits. The skin biopsy and direct immune fluorescent examination (DIF) were not performed during that time. Topical steroids and oral hydroxychloroquine 200mg twice a day were prescribed. After 9 years of irregular follow up, this patient was found to have nasopharyngeal cancer and received radiotherapy. A repeated cutaneous physical examination revealed no Heliotrope sign, Gottron papules, perungual erythema, poikiloderma, or shawl sign. No muscular weakness was found either. The repeated laboratory examination for autoantibody confirmed positive anti-Ro and anti-La antibodies, and negative anti-Sm/RNP antibody. Six months later, he developed muscular weakness and elevated creatine phosphokinase (CPK: 2,266 IU/L). He was then diagnosed as dermatomyositis. After 1 year, multiple persistent itching round infiltrated erythematous plaques developed on the upper back and bilateral arms (Figure 1 and 2). A skin biopsy was performed on the elevated plaque of the back and the sample was sent for histological examination. Microscopic examination showed perivascular interstitial mixed-cell infiltrate, composed of neutrophils, histiocytes, lymphocytes, and nuclear dusts within the superficial and deep dermis (Figure 3 and 4). In some foci, the palisaded granuloma was...
found without vasculitis, features consistent with palisaded and neutrophilic granulomatous dermatitis. The absence of palisaded granulomatous inflammation with central necrobiosis and mucin deposition ruled out the possibility of granuloma annulare. The DIF of the skin biopsy showed only weak granular IgG and linear IgM deposition along dermo-epidermal junction, without definite diagnostic conclusion. The skin lesions improved after treatment with topical clobetasol propionate ointment.

Discussion

Palisaded and Neutrophilic Granulomatous Dermatitis (PNGD) is a rare cutaneous manifestation, which is most commonly reported with rheumatoid arthritis [1]; however, it has also been associated with systemic lupus erythematosus [2-9], inflammatory bowel disease [10], and systemic sclerosis [11]. Other related diseases like Behcet disease, hepatitis, sarcoidosis, leukocytoclastic vasculitis has been reported, too [12-15]. To the best of our knowledge, PNGD associated with dermatomyositis has never been reported so far [16]. The clinical manifestations of PNGD include asymptomatic or intensely painful papules, nodules, linear subcutaneous in durated cordlike bands (the burning rope sign) [4], and plaques on multiple body sites. The histological examination typically shows a dense neutrophilic infiltrate with degenerated collagen, leukocytoclastic debris, and palisading granulomas, without vasculitis [3]. Chu et al proposed that the histological appearances of palisaded neutrophilic granulomatous dermatitis might vary from early (dense inflammatory infiltrates, composed of lymphocytes, neutrophils, histiocytes, and eosinophils) to late stages (palisading granulomas with fibrosis) of the disease [3]. The clinical and pathological presentations of our patient correlate with the previous findings in the literature.

The etiology of PNGD is unknown. Finan and Winkelmann observed IgM and C3 in small vessels by using DIF in 1983 [17]. They presumed that the cutaneous lesions were the result of immune complexes generated by underlying systemic diseases. Although the DIF of this patient did not show any deposition along the small vessels, there were still weak IgG and IgM depositions along dermo-epidermal junction, which might be due to the underlying dermatomyositis. The neutrophilic infiltrate in dermatomyositis related PNGD may be explained by the deposition of the immune complex and further activation of complement cascade. The adaptive immune system, particularly through the T helper lymphocytes (Th) 17-cell subset, may also secrete neutrophil-recruiting chemokines, such as IL-17A and G-CSF. Local synthesis of IL-17A by infiltrating Th17 cells in skin lesions of Autoimmune Connective Tissue Diseases (AICTDs) could lead to the presence of a nonspecific neutrophilic infiltrate in PNGD of AICTDs [18]. Neutrophils may also be over recruited in AICTD skin lesions because of an abnormal expression and/or activation of adhesion/migration molecules through the endothelial barrier of inflamed tissues [19]. Caproni et al proposed that the neutrophilic infiltrate described in specific skin lesions of dermatomyositis could be explained by increased expression of adhesion molecules, such as vascular-cell adhesion molecule-1, intercellular adhesion molecule-1, and E-selections on endothelial cells [20]. Therefore, it is possible that the deposition of immune complexes in dermal vessels from underlying autoimmune disease activates the complement and Th17 lymphocytes, leading to over recruited aggregation of neutrophils and other inflammatory cells, causing degeneration and ischemia of collagen, followed by granulomatous reaction to this damaged collagen. According to Rosenbach and English [16], the unifying term “reactive granulomatous dermatitis” was proposed to encompass PNGD, interstitial granulomatous dermatitis, and interstitial granulomatous drug reaction, in order to emphasize these specific cutaneous granulomatous reaction patterns that occur in the setting of a systemic trigger.

In conclusion, we reported a rare association between PNGD and dermatomyositis. A new term, reactive granulomatous dermatitis, is suggested to describe these diverse clinical and pathological reaction patterns in response to some underlying systemic process.

References


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